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TRANSMITTAL		Filing Date	July 9, 2003				
FORM		First Named Inventor	Bizzarro				
(to be used for all correspondence after initial filing)		Art Unit	1626				
		Examiner Name	Wright, Sonya N				
Total Number of Pages in This Submission	al Number of Pages in This Submission			20400 US6			
ENCLOSURES (Check all that apply)							
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Date March 4, 2004							
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I hereby certify that this correspondence is being transmitted to the USPTO or deposited with the United States Postal Service with sufficient postage as first class mail in an envelope addressed to: Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313 on this dat March 4, 2004							
Typed or printed Eileen M. Ebel							
Signature Sil, MAS					Date	March 4,	2004

This collection of information is required by 37 CFR 1.5. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.14. This collection is estimated to take 12 minutes to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, Washington, DC 20231. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.



PATENT APPLICATION

Group: 1626

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Patent Application

Fred Thomas Bizzarro, et. al.

Serial No. 10/616,359, filed July 9, 2003

Docket No.: 20400 US6

For: HETEROAROMATIC GLUCOKINASE ACTIVATORS

RESPONSE UNDER 35 U S C § 121

Nutley, New Jersey 07110 March 4, 2004

Examiner: Wright, Sonya N

Commissioner for Patents P.O. Box 1450 Alexandria, VA 22313-1450

Dear Sir:

In response to the Office Action dated February 6, 2004, Applicants provide the following remarks.

In response to the requirement for restriction, Applicants elect Group II, claims 1-149 and 267-278, directed to compounds of formula I wherein R⁴ is thiazole and thiadiazole.

In terms of a species election, Applicants request that the Examiner begin the search for compounds in which R² is methylsulfonyl, i.e., SO₂CH₃. Applicants identify species Example numbers 3, 12, 13, 89 and 97. In these Examples, R³

Serial No. 10/616,359 Filed: July 9, 2003

is cyclopentyl. In Examples 3 and 13, R^1 is hydrogen. In Examples 12 and 97, R^1 is CI. In Example 89, R^1 is CF_3 .

Applicants traverse the requirement for restriction, however.

The U.S. Patent Office is required to examine on the merits the entirety of generic claims. Applicants point out that it is well-established law that restriction within a single claim cannot be sustained under 35 U.S.C. §121. As is stated in *In re Weber*, 198 USPQ 328 (CCPA 1978), at pages 331-332:

"§121 provides the Commissioner with the authority to promulgate rules designed to *restrict an application* to one of several claimed inventions when those inventions are found to be "independent and distinct." It does not, however, provide a basis for an examiner acting under the authority of the Commissioner to *reject* a particular *claim* on that same basis." (Emphasis in original text.)

If the Patent Office were to withdraw applicants' claims in part from further consideration due to an intra-claim restriction, the requirement amounts in fact to a rejection, see *In re Hass*, 179 USPQ 623, 625 (CCPA 1973).

Applicants have the right under U.S. patent law to claim their invention using the limitations that they regard as essential to delineate the invention, as long as the requirements of 35 U.S.C. §112 are met.

The law does not authorize the U.S. Patent Office to derive its own concept of a generic form of the claimed subject matter and require that it be carved out of existing claims. Applicants have the right under U.S. patent law to claim their invention using the limitations they provide to delineate the invention, as long as the requirements of 35 USC § 112 are met. See *In re Weber* at 331 and *In re Wolfrum and Gold*, 179 USPQ 620, 622 (CCPA 1973).

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Also, the Examiner is required to follow the U.S. Patent Office procedures set forth in MPEP § 809. The Examiner must perform a search for all claims readable on the elected species. The elected species should be examined. If the species is patentable, then the next species should be examined, and so forth, until an unpatentable species is found. If no species is found unpatentable, then the generic claim should be allowed.

Applicants request that the intra-claim restriction requirement be withdrawn.

It is believed that no fees are due. If a fee is owing, please charge our deposit account no. 08-2525

Respectfully submitted,

Attorney for Applicant(s)

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